

Challenges in dengue research: A computational perspective

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Abstract

The dengue virus is now the most widespread arbovirus affecting human populations, causing significant economic and social impact in South America and South-East Asia. Increasing urbanization and globalization, coupled with insufficient resources for control, misguided policies or lack of political will, and expansion of its mosquito vectors are some of the reasons why interventions have so far failed to curb this major public health problem. Computational approaches have elucidated on dengue's population dynamics with the aim to provide not only a better understanding of the evolution and epidemiology of the virus but also robust intervention strategies. It is clear, however, that these have been insufficient to address key aspects of dengue's biology, many of which will play a crucial role for the success of future control programmes, including vaccination. Within a multiscale perspective on this biological system, with the aim of linking evolutionary, ecological and epidemiological thinking, as well as to expand on classic modelling assumptions, we here propose, discuss and exemplify a few major computational avenues—real-time computational analysis of genetic data, phylodynamic modelling frameworks, within-host model frameworks and GPU-accelerated computing. We argue that these emerging approaches should offer valuable research opportunities over the coming years, as previously applied and demonstrated in the context of other pathogens.

KEYWORDS

computation, dengue, epidemiology, evolution, models

1 | BACKGROUND

1.1 | The dengue virus

Dengue virus (DENV) is a single-stranded, positive-sense RNA virus with a population comprised of four antigenically distinct groups, or serotypes (DENV1–4), that present similar population dynamics and are the cause of clinically indistinguishable illnesses in humans (Gubler, 1998; Halstead, 2007; Kyle & Harris, 2008). Classification based on genomic organization, homology and antigenic cross-reactivity place DENV into the *Flavivirus* genus of the viral family *Flaviviridae* (Calisher,

Karabatsos, Dalrymple, Shope, & Porterfield, 1989). More than 70 flaviviruses have so far been identified (Kuno, Chang, Tsuchiya, Karabatsos, & Cropp, 1998), many of which are important human pathogens, including the hepatitis C virus (HCV), Japanese encephalitis virus (JEV), West Nile virus (WNV), yellow fever virus (YFV) and the recently emerged Zika virus (ZIKV). Among the ones that are vector-borne (all except HCV), DENV is responsible for the highest human morbidity and mortality, with children carrying the highest burden of disease (Anders et al., 2011; Jain & Chaturvedi, 2010; Murray, Quam, & Wilder-Smith, 2013).

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1.2 | Dengue evolution

Observations were made in the late 1970s of dengue-like viruses and respective animal reservoirs in Asia and West Africa (Aiken & Leigh, 1978). Since then, a wide number of sylvatic strains (i.e., of animal origin) have been isolated and reported to belong to each of the lineages of human-bearing serotypes, proposing a zoonotic origin for the pathogen (Holmes & Twiddy, 2003). Introduction of each serotype into the human population was extrapolated to have taken place at different time points, with DENV2's most recent common ancestor being the oldest, estimated to be approximately 1,000 years old (Twiddy, Holmes, & Rambaut, 2003; Weaver & Vasilakis, 2009), roughly corresponding to some of the first reports of dengue-like illnesses (Vasilakis & Weaver, 2008; Tang et al., 2010; Weaver & Vasilakis, 2009). The genetic distance between currently circulating serotypes is likely a by-product of allopatric evolution due to ecological partitioning of ancestral sylvatic strains (Holmes & Twiddy, 2003; Weaver & Vasilakis, 2009). This hypothesis is sustained not only by DENV's current population structure, but also by studies that report sylvatic and endemic (i.e., human bearing) strains to exhibit similar evolutionary rates and (cell-) infection potential (Twiddy et al., 2002; Vasilakis et al., 2007).

Within each serotype, nucleic acid sequencing techniques have identified genetic subgroups or lineages, commonly denominated as dengue genotypes (Tang et al., 2010; Weaver & Vasilakis, 2009). Notably, the overall amino acid sequence similarity among genotypes is generally above 90% (Holmes & Twiddy, 2003; Leitmeyer et al., 1999), whereas between serotypes, it can range between 60% and 80% (Irie, Mohan, Sasaguri, Putnak, & Padmanabhan, 1989; Rothman, 2011). Although RNA viruses generally present rapid evolution and accumulation of diversity occurring at a similar time-scale as their host's ecological dynamics (Grenfell, Pybus, Gog, Wood, & Daly, 2004), studies have demonstrated that strong purifying selection dominates DENV's recent evolution (Bennett, Holmes, Chirivella, Rodriguez, & Beltran, 2003; Holmes, 2003; Holmes & Twiddy, 2003; Weaver & Vasilakis, 2009; Zhang, Mammen, Chinnawirotpisan, Klungthong, & Rodpradit, 2005). A prevailing hypothesis is that the virus has evolved to replicate efficiently in both the vertebrate and arthropod hosts (Filomatori, Carballeda, Villordo, Aguirre, & Pallarés, 2017), expressing a compromise genome in which most amino acid changes are expected to be deleterious and selectively removed from the population (Holmes, 2003; Holmes & Twiddy, 2003; Weaver & Vasilakis, 2009). The lack of recombination, supported by rare reports on naturally occurring recombinants (Waman, Kolekar, Ramtirthkar, Kale, & Kulkarni-Kale, 2016; Weaver & Vasilakis, 2009), is further believed to limit the fixation and emergence of single, positively selected mutations due to clonal interference (Grenfell et al., 2004; Lourenço & Recker, 2010; Miralles, Gerrish, Moya, & Elena, 1999).

1.3 | Dengue transmission cycle

Dengue virus is transmitted to humans by the bite of infectious mosquitoes. Two mosquito species have been identified as the main disease

vectors: *Aedes aegypti* and *Ae. albopictus*. Intercontinental changes in the second half of the 20th century, in particular the post-World War II boom in urbanization and globalization, have facilitated the introduction and establishment of dengue's two principal mosquito vectors into many major urban and peri-urban settings and resulted in the worldwide establishment of endemic transmission cycles (i.e., between humans; Vasilakis, & Weaver, 2008; Murray et al., 2013; Weaver & Vasilakis, 2009). *Ae. aegypti* is particularly domesticated and often uses artificial water containers found within and around human habitats as breeding sites. Adult mosquitoes are optional haematophages, but females require blood for egg production and show strong preference for human blood meals (Scott, Morrison, Lorenz, Clark, & Strickman, 2000).

In contrast, the transmission cycle of sylvatic viruses takes place in forest environments of South-East Asia and West Africa between nonhuman primates and arboreal *Aedes* species. Rural populations living in close proximity to forested or plantation areas are periodically exposed to such variants (Holmes & Twiddy, 2003; Vasilakis, & Weaver, 2008; Weaver & Vasilakis, 2009). Secondary transmission is rarely reported, which may be a combined result of limited human-to-vector transmission potential, underreporting due to low pathogenicity, weak surveillance and lack of peridomestic vectors (*Ae. aegypti* and *Ae. albopictus*).

1.4 | Dengue epidemiology

The epidemiological picture of dengue across the world is variable. While hyperendemicity is classically associated with South-East Asia, the 21st century has witnessed the endemic establishment of the virus in South and Central America (Brathwaite Dick, San Martín, Montoya, del Diego, & Zambrano, 2012; San Martín, Brathwaite, Zambrano, Solórzano, & Bouckennooghe, 2010). Dengue has also recently emerged in the United States, with repeated but so far self-contained outbreaks in Florida and Texas (Guzman, Halstead, Artsob, Buchy, & Farrar, 2010; Radke, Gregory, Kintziger, Sauber-Schatz, & Hunsperger, 2012; Thomas, Santiago, Abeyta, Hinojosa, & Torres-Velasquez, 2016). In Europe, outbreaks in Italy, France, Greece and the Madeira island (Portugal) have been reported in recent years, but with no observations of endemic circulation (Cavirini, Gaibani, Pierro, Rossini, & Landini, 2009; Gautret, Botelho-Nevers, Charrel, & Parola, 2010; Gjenero-Margan, Aleraj, Krajcar, Lesnikar, & Klobučar, 2011; Lourenço & Recker, 2014).

Poor surveillance, no official reporting to the WHO (Jaenisch, Junghanss, Wills, Brady, & Eckerle, 2014) and the high prevalence of other diseases makes the situation in Africa more uncertain. Recently, outbreaks of dengue or dengue-like illness were reported in dry countries of Eastern Africa and South West Asia, such as Saudi Arabia (Aziz, Al-Shami, Mahyoub, Hatabbi, & Ahmad, 2014), Yemen (Bin Ghouth, Amarasinghe, & Letson, 2012), Sudan (Seidahmed, Hassan, Soghaier, Siam, & Ahmed, 2012) and Madagascar (Schwarz, Girmann, Randriamampionona, Bialonski, & Maus, 2012). In 2004 and 2009, epidemics occurred in the isolated populations of Cape Verde (Franco, Di Caro, Carletti, Vapalahti, & Renaudat, 2010), Mauritius islands (Issack, 2010) and Reunion island (Larrieu,

Dehecq, Balleydier, Jaffar, & Michault, 2012), and a major epidemic in Pakistan took place in 2013 (Kraemer, Perkins, Cummings, Zakar, & Hay, 2015; Khan, Khan, & Amin, 2016). Together with the widespread presence of *Ae. aegypti* and *Ae. albopictus*, this would indicate that the burden of dengue in Africa is likely to be worse than previously acknowledged (Guzman et al., 2010; Jaenisch et al., 2014; Kyle & Harris, 2008).

Driven by seasonal variation in vector abundances, incidence patterns are usually characterized by annual outbreaks (Figure 1a; Cuong, Hien, Duong, Phong, & Cam, 2011; Lourenço & Recker, 2013; Nisalak, Endy, Nimmannitya, Kalayanaroj, & Thisayakorn, 2003). In addition, multi-annual epidemics with a 2- to 5-year period have also been reported in highly endemic settings (Figure 1a,b; Bennett, Drummond, Kapan, Suchard, & Muñoz-Jordán, 2010; Cummings, Irizarry, Huang, Endy, & Nisalak, 2004; Cuong et al., 2011; Lourenço & Recker, 2013; Nisalak et al., 2003; Thai, Cazelles, Nguyen, Vo, & Boni, 2010). In contrast, regions where transmission has been historically low or where dengue serotypes were only recently introduced are more likely to show wave-like, increasing patterns of incidence (Figure 1c,f). Spatial heterogeneities are also common in reported data, highlighting that the success of dengue viruses can be dictated by numerous demographic and ecological determinants (Figure 1f; Adams & Kapan, 2009; Beebe, Cooper, Mottram, & Sweeney, 2009; Campbell, Haldeman, Lehnig, Munayco, & Halsey, 2015; Chew, Woon, Amin, Adnan, & Abdul Wahab, 2016; Cummings et al., 2004; Lourenço & Recker, 2010; Lourenço & Recker, 2013).

Where all four serotypes cocirculate endemically, relative prevalence levels of each serotype are characterized by oscillatory behaviour

with periods of 8–10 years (Figure 1d,e; Adams, Holmes, Zhang, Mammen, & Nimmannitya, 2006; Bennett et al., 2010; Lourenço & Recker, 2013; Nisalak et al., 2003). Whereas incidence dynamics are thought to be driven by a complex interplay of local factors, from the abundance of susceptible humans, climate and viral traits, to the density and distribution of both the human and the vector populations (Johansson, Dominici, & Glass, 2009; Morrison, Minnick, Rocha, Forshey, & Stoddard, 2010; Nagao, Svasti, Tawatsin, & Thavara, 2008; Tsuzuki, Vu, Higa, Nguyen, & Takagi, 2009; Williams, Bader, Kearney, Ritchie, & Russell, 2010), there is lack of evidence that these factors directly influence the qualitative properties of serotype cyclical behaviour, which seems to be a universal feature of dengue epidemiology.

1.5 | Human infections, immunity and disease

Primary infections in humans with any of the four serotypes are generally mild with the majority of cases believed to be asymptomatic (Kyle & Harris, 2008; Nisalak et al., 2003). Infections usually result in a short-lived, febrile illness, which is commonly referred to as *dengue fever* (DF) and is characterized by arthralgia (joint pain), myalgia (muscle pain), low-grade fever and retro-orbital headaches (Balmaseda, Hammond, Tellez, Imhoff, & Rodriguez, 2006; Burke, Nisalak, Johnson, & Scott, 1988; Wichmann, Yoon, Vong, Limkittikul, & Gibbons, 2011). In some cases, this may progress to more severe and life-threatening forms of disease—*dengue haemorrhagic fever* (DHF) or *dengue shock syndrome* (DSS)—with manifestations of circulatory failure, vascular permeability and haemorrhagic symptoms (Gubler, 1998; Halstead, 2007).

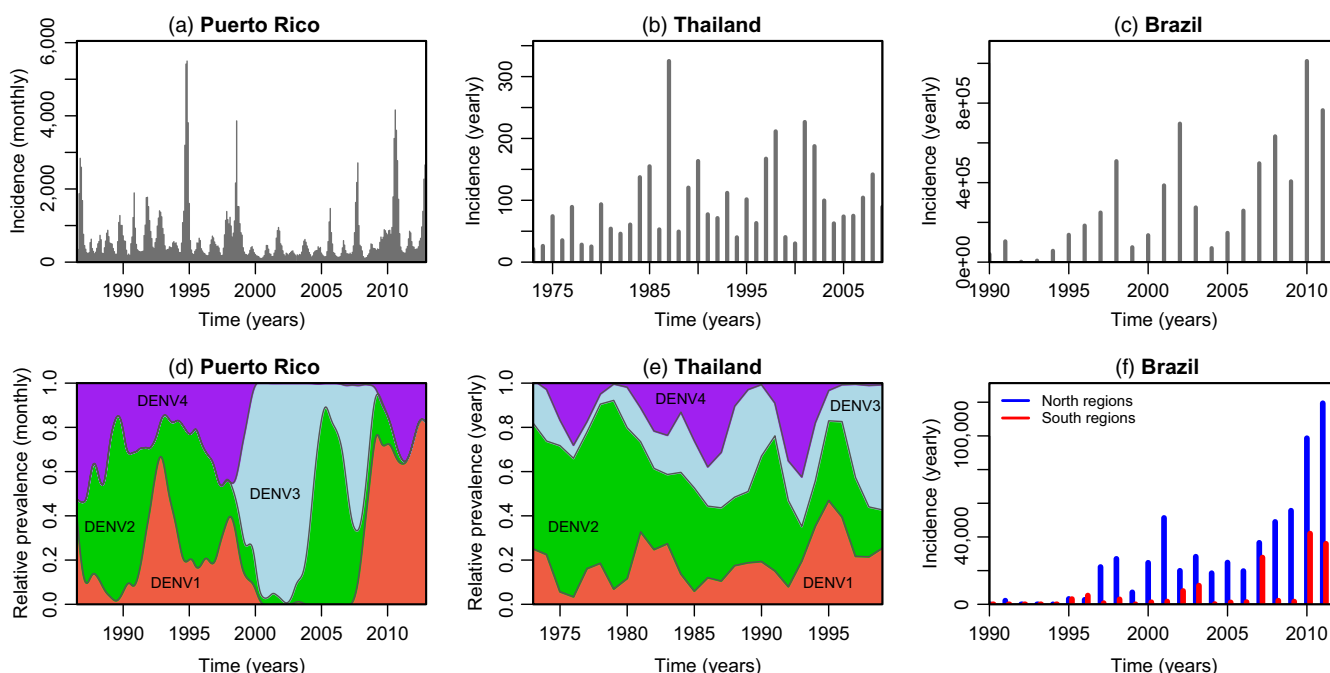


FIGURE 1 Epidemiological time series of reported dengue cases for three different countries. (a) Total monthly incidence for Puerto Rico. (b) Annual dengue haemorrhagic fever incidence per 100,000 individuals in Thailand. (c) Total annual incidence for Brazil. (d) Monthly serotype, relative prevalence for Puerto Rico. (e) Yearly serotype, relative prevalence for Thailand. (f) Yearly absolute incidence for the north and south of Brazil (see Appendix: Data Description for details)

Flaviviruses are known to induce the phenomenon of *original antigenic sin* (OAS) (Francis, 1960), in which B-cell clones sequentially challenged by viruses of the same family initially respond by synthesizing antibodies with higher affinity to a previous infecting virus (Halstead, Rojanasuphot, & Sangkawibha, 1983). This creates a cascade of regulatory changes that facilitates expression and synthesis of pro-inflammatory cytokines and induces a positive feedback that increases the rate of intracellular infection, leading to higher viraemia levels and worse clinical outcomes (Beltramello, Williams, Simmons, Macagno, & Simonelli, 2010; Fernandez-Garcia, Mazzon, Jacobs, & Amara, 2009; Green, Vaughn, Kalayanaraj, Nimmannitya, & Suntayakorn, 1999; Ubol & Halstead, 2010; Whitehorn & Simmons, 2011). In dengue infections, this cascade is commonly referred to as *antibody-dependent enhancement* (ADE) and is mainly associated with heterologous DENV challenges (Balsitis, Williams, Lachica, Flores, & Kyle, 2010; Halstead, Mahalingam, Marovich, Ubol, & Mosser, 2010; Wrammert, Onlamoon, Akondy, Perng, & Polsrila, 2012) or the presence of maternal antibodies in infants (Chau, Hieu, Anders, Wolbers, & Lien, 2009).

Generally, cellular immune responses (T cell) to DENV are seen to cover most of the viral proteins (Duangchinda, Dejnirattisai, Vasanawathana, Limpitkul, & Tangthawornchaikul, 2010; Rothman, 2011), but with a particular dominant signal for NS3 (Duangchinda et al., 2010), which represents only $\approx 20\%$ of DENV amino acid coding sequence (Rothman, 2011). T-cell clones can be serotype cross-reactive, sero-specific or even strain-specific (i.e., within serotype; Bashyam, Green, & Rothman, 2006; Mangada, Endy, Nisalak, Chunsuttiwat, & Vaughn, 2002; Rothman, 2011; Sierra, García, Pérez, Morier, & Rodríguez, 2002; Ubol & Halstead, 2010). The cross-reactivity is not surprising, however, given the $\approx 70\%$ amino acid identity between DENV1 and DENV4 (Rothman, 2011). OAS in T-cell responses, including inflammation and cytokine-induced alteration of vascular permeability (Rothman, 2011), have also been described in secondary infections (Mongkolsapaya, Dejnirattisai, Xiao-ning, Vasanawathana, & Tangthawornchaikul, 2003), with consequences for pathogenicity (Bashyam et al., 2006; Duangchinda et al., 2010; Imrie, Meeks, Gurary, Sukhbataar, & Kitsutani, 2007; Mangada et al., 2002; Stephens, Klaythong, Sirikong, Vaughn, & Green, 2002).

Hence, available evidence from both humoral and cellular immunity suggests that the clinical outcome of DENV infection is finely tuned to an individual's immune response, in particular to the degree of OAS and associated inflammatory levels. Factors that may influence the regulation of an individual's response, in particular for secondary challenges, might include the following: host genetics, viral strain, sequence of infecting serotypes, age, co-infection with other pathogens, etc. However, there is a major gap in our current knowledge on reliable, immunological markers for clinical outcomes, and discerning between such factors is often difficult (Rothman, 2011), although, generally, the most significant risk factor for severe disease is secondary infection with an heterologous serotype as a primer for OAS (Burke et al., 1988; Guzman & Kouri, 2008; Halstead et al., 2010; Kyle & Harris, 2008; Sangkawibha, Rojanasuphot, Ahandrik, Viriyapongse, & Jatanasen, 1984; Thein, Aung, Shwe, Aye, & Zaw, 1997; Rothman &

Ennis, 1999; Halstead & O'Rourke, 1977; Dejnirattisai, Jumnainsong, Onsirakul, Fitton, & Vasanawathana, 2010).

Recovery is thought to result in lifelong protection against the infecting serotype (homologous immunity) (but see (Forshey, Reiner, Olkowski, Morrison, & Espinoza, 2016; Waggoner, Balmaseda, Gresh, Sahoo, & Montoya, 2016) for possible incomplete protection). A period of 6–36 months of temporary cross-protective immunity to all serotypes has also been suggested after recovery (see e.g., Sabin, 1952; Reich, Shrestha, King, Rohani, & Lessler, 2013), although it is still uncertain whether this form of immunity solely protects against clinical disease or completely prevents infection (Bhoomiboonchoo, Nisalak, Chansatiporn, Yoon, & Kalayanaraj, 2015; Anderson et al., 2014; Kochel, Watts, Halstead, Hayes, & Espinoza, 2002; Kochel, Watts, Gozalo, Ewing, & Porter, 2005; Olkowski, Forshey, Morrison, Rocha, & Vilcarromero, 2013). Upon recovery from a secondary, heterologous infection, humans are believed to be (clinically) protected against further challenges. Due to the rarity of third and fourth infections in clinical data and the difficulties in assessing an individual's infection history because of high antibody cross-reactivity in serological assays (Halstead, 2007; Nisalak et al., 2003; Olkowski et al., 2013; Gibbons, Kalanaraj, Jarman, Nisalak, & Vaughn, 2007), it is not yet clear to what degree secondary infection prevents re-infections or simply renders subsequent infections, even by previously un-experienced serotypes, clinically silent.

1.6 | Immune interactions and computational approaches

Mathematical and computational models are recognized as an important tool in infectious disease epidemiology and have been applied to dengue since the mid-20th century. Hand-in-hand with the increase in the global distribution and burden of dengue, the number of modelling studies has also gone up (Figure 2). Topics have ranged widely, from estimations of dengue's reproductive number to the impact of climate and space on disease transmission and spread, to the evaluation and assessment of disease control strategies. Dengue's complex epidemiological dynamics, with multi-annual outbreaks and sequential replacement of dominant serotypes, has also attracted a fair share of the attention of the modelling community, with many studies concentrating on immune interactions between dengue serotypes, either through ADE and/or by means of cross-immunity. In fact, the incorporation of immune interaction has become ubiquitous to most dengue models and hence deserves a little more detail here.

It is broadly recognized that immune competition can act to desynchronize antigenically diverse pathogens, such as dengue, malaria (*Plasmodium falciparum*) or the influenza A virus, and cause nonlinear or chaotic oscillations not too dissimilar to the observed epidemiological patterns of dengue. For dengue, competition between serotypes can be driven by ADE of secondary, heterologous infections or through (temporary or permanent) cross-immunity. Both types of interactions are assumed to affect the transmissibility of subsequent infections, leading to frequency-dependent selection that has been demonstrated to generate oscillatory behaviour and further facilitate serotype persistence (Schwartz, Shaw, Cummings, Billings, & McCrary,

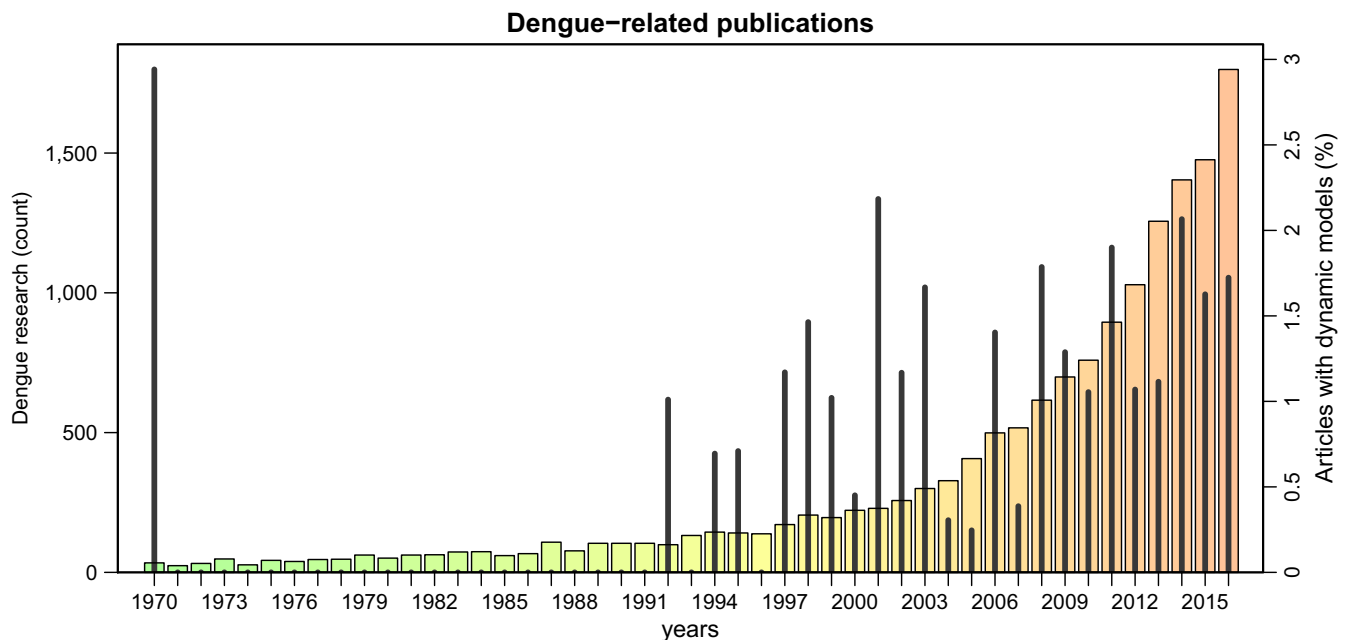


FIGURE 2 Dengue publication over the last five decades. Total number of dengue articles per year (bars) and the percentage of those with a computational focus (spikes). Between 1970 and 2016, a total of 15,267 dengue articles were published, including 190 modelling studies (see Appendix for details)

2005; Ferguson, Anderson, & Gupta, 1999; Cummings, Schwartz, Billings, Shaw, & Burke, 2005; Aguiar, Kooi, & Stollenwerk, 2008; Wearing & Rohani, 2006; Adams & Boots, 2006; Hu, Thoens, Bianco, Edlund, & Davis, 2013).

With regard to ADE, initial criticism concerning the level of enhancement required for serotype desynchronization could be countered by also assuming enhancement in the susceptibility to secondary infections (see e.g., (Recker, Blyuss, Simmons, Hien, & Wills, 2009)). Importantly, however, empirical evidence supporting a universal increase in the transmission potential of secondary infections is still lacking. That is, while the ADE phenomenon has been shown both in *in vitro* and *in vivo* studies (Halstead et al., 2010; Dejnirattisai et al., 2010; Halstead, 2011), its actual onsequences for disease transmission and the proportion of hosts contributing to ADE-induced phenotypes at any given time are largely unknown.

Cross-immunity, where an infection by one serotype negatively affects the fitness of subsequently infecting serotypes, has equally been put forward in modelling studies to explain dengue's complex epidemiology (Adams et al., 2006; Reich et al., 2013; Aguiar et al., 2008; Wearing & Rohani, 2006). Recent studies on prospective cohorts have found that shorter time periods between sequential infections are associated with some degree of clinical protection (Anderson et al., 2014; Montoya, Gresh, Mercado, Williams, & Vargas, 2013). However, apart from a single human study conducted in the 1950s (Sabin, 1952), experimental support for both the nature and duration of cross-immunity is still lacking. Consequently, there is no broad consensus about the protective efficacy of the observed cross-reactivity between dengue serotypes and whether cross-reactive responses prevent infection or solely act to alleviate clinical outcomes. Model assumptions have therefore diverged

significantly with regard to the duration and the type of protection following a primary dengue infection.

It is worth noting that some of the controversies surrounding serotype immune interactions and their effects on the epidemiological dynamics of dengue are partially consequences of the actual modelling frameworks themselves. That is, deterministic models based on mass-action principles, which comprise the majority of mathematical models found in the published literature, require an interaction term in order to desynchronize the serotypes and to exhibit dengue's characteristic multi-annual incidence patterns. Stochastic, and in particular individual-based, models do not rely on this assumption to generate epidemiological time series with such properties. The contrasting dynamic behaviour is driven by the amplification of stochastic effects at the individual-level (demographic stochasticity), which keeps each serotype/strain in a transient regime rather than approaching the expected deterministic equilibrium even in the absence of explicit desynchronizing factors (Lourenço & Recker, 2013; Alonso, McKane, & Pascual, 2007). Crucially, understanding the epidemiological consequences of dengue immune interactions goes beyond the theoretical argument and has recently been shown to be of major importance for dengue vaccination impact studies based on mathematical models (Lourenço & Recker, 2016; Flasche et al., 2016).

2 | CHALLENGES AND OPPORTUNITIES IN COMPUTATIONAL RESEARCH

The zoonotic origin, separate endemic and sylvatic transmission cycles, immunological serotype interactions and the vast diversity of

dengue genotypes provide a valuable background for integrative evolutionary approaches in the context of dengue. However, the predominant focus on single biological scales (e.g., host-host transmission) has so far limited our capacity to link important population genetic and ecological observations with the pathogen's epidemiological behaviour within a single framework. In sharp contrast to other major viral pathogens, such as the influenza A virus or the human immunodeficiency virus (HIV), evolutionary frameworks do not feature prominently in dengue's modelling literature, with the field of phylogenetics so far dominating evolutionary approaches.

In general, the continuing geographical and epidemic expansion of dengue's genetic variants and mosquito vectors, and the failure to prevent the introduction and establishment of sustained transmission across the globe, call for a renewed focus and advancement in computational approaches. In the context of dynamic modelling, the literature has been dominated by frameworks based on ordinary differential equations (ODE), which are tractable and computationally inexpensive. However, such approaches often take key assumptions unquestioningly, such as homogeneous mixing, competition factors to desynchronize antigenic types, or continuity and determinism, which can hamper our understanding of the major determinants of dengue's evolution, epidemiology and control. Within a multiscale perspective on this biological system, and with the aim of linking evolutionary, ecological and epidemiological aspects as well as to expand on classic modelling assumptions, we here propose, discuss and exemplify a few major computational avenues, which we hope should offer valuable research opportunities over the coming years: real-time computational analysis of genetic data, phylodynamic modelling frameworks, within-host model frameworks and GPU-accelerated computing.

2.1 | Real-time collection and computational analysis of genetic data

Like the yellow fever virus (Auguste, Lemey, Pybus, Suchard, & Salas, 2010), DENV was probably first introduced in the Americas through infected hosts and vectors during the slave trade (Weaver & Vasilakis, 2009; Vasilakis, & Weaver, 2008). At present, the distribution of DENV serotypes results from a combination of strong population structure and gene flow across geographical regions. South-East Asia harbours the greatest DENV genetic diversity, and it is from there that viral lineages seem to seed epidemics in the Americas and elsewhere (Holmes, 2008). Several studies suggest that every 7–10 years, DENV genotypes become extinct and the replacement by new viral lineages leads to period fluctuations in genetic diversity (Klungthong, Zhang, Mammen, Ubol, & Holmes, 2004; Thu, Lowry, Jiang, Hlaing, & Holmes, 2005; Nunes, Palacios, Faria, Sousa, & Pantoja, 2014). Each of these epidemics is fuelled by a combination of local herd susceptibility (Salje, Lessler, Endy, Curriero, & Gibbons, 2012), seasonal abundance of the *Ae. aegypti* mosquito (Kraemer, Sinka, Duda, Mylne, & Shearer, 2015) and an increasingly well-connected mobility network that facilitates the movement of infected hosts and vectors (Nunes et al., 2014; Wesolowski, Qureshi, Boni, Sundsøy, & Johansson, 2015).

Dengue virus incidence is often underestimated and novel surveillance, tools need to be applied to improve current estimates and reduce misclassification (Silva, Rodrigues, Paploski, Kikuti, & Kasper, 2016). The most commonly used dengue diagnostic tests are based on antibody detection and often show cross-reactivity with other flaviviruses, including West Nile virus, Japanese encephalitis virus and yellow fever virus. Classification based on clinical symptoms alone is complicated by the circulation of pathogens that cause similar clinical symptoms to DENV infection, such as the chikungunya virus (CHIKV) and ZIKV, and may thus hamper accurate descriptions of DENV dynamics (Faria, Lourenço, Marques de Cerqueira, Maia de Lima, & Carlos Junior Alcantara, 2016; Roth, Mercier, Lepers, Hoy, & Duituturaga, 2014).

One of the ultimate challenges in infectious disease control is to predict and halt the course of an epidemic following its emergence. However, insufficient surveillance can obfuscate predictions of virus dynamics across local, regional and global scales. For instance, diagnostic tools and reporting are unevenly supported across locations with risk for transmission. This is particularly the case for central Africa, for which little information is known about the expansion of arboviruses (Jaenisch et al., 2014), although molecular detection from returning travellers has revealed active undetected transmission of DENV and CHIKV in this region (Parreira, Centeno-Lima, Lopes, Portugal-Calisto, & Constantino, 2014) and has identified it as the source of the CHIKV East-Central-South-African genotype in the American continents (Nunes, Faria, de Vasconcelos, Golding, & Kraemer, 2015). It is also the case that representative data are essential to quantify the burden of transmission and to predict dynamic behaviour and impact of interventions (Jaenisch et al., 2014). However, even in regions with access to diagnosis and surveillance, it is rarely the case that reported case data on arboviruses are representative (Silva et al., 2016; Lourenço, Maia de Lima, Faria, Walker, & Kraemer, 2017). A major example for which both of these issues are particularly challenging is the Amazon region, that has been suggested to play a stepping stone role in the introduction of DENV lineages from the Caribbean into Brazil (Temporão, Penna, Carmo, Coelho, & Azevedo, 2011; Nunes, Faria, Vasconcelos, de Almeida Medeiros, & de Lima, 2012), and may have also been involved in the emergence and maintenance of ZIKV (Faria, Quick, Claro, Thézé, & de Jesus, 2017).

With the ongoing democratization of pathogen sequencing (Faria, Sabino, Nunes, Alcantara, & Loman, 2016), it is hoped that the generation of genetic data will soon become standard in public health and clinical practise (Houldcroft, Beale, & Breuer, 2017). Public databases are now becoming populated with thousands of partial and complete genome sequences. Sequencing during epidemics is even becoming an increasing trend, which is important for our understanding on how particular genotypes associated with increased virulence may be generated and eventually fixate (OhAinle, Balmaseda, Macalalad, Tellez, & Zody, 2011). Moreover, genetic data collected over the course of months, years and decades allow for a better understanding of the dynamics of DENV lineages at the population level, offering key insights into demographic, ecological and genetic factors which are difficult, if not impossible, to track with clinical records alone.

With the *real-time genomic epidemiology revolution* (Gardy, Loman, & Rambaut, 2015), new and complementing insights (to clinical reported case data) are expected to provide unique information about the macro- and microscale process shaping viral dynamics. Statistical models that take into account shared ancestry allow testing for the contribution of demographic factors associated with virus persistence and spread (Faria, Suchard, Rambaut, Streicker, & Lemey, 2013; Lemey, Rambaut, Bedford, Faria, & Bielejec, 2014). Using such approaches, air travel has been shown to play a pivotal role in the spread of vectors (Tatem, Hay, & Rogers, 2006) and DENV (Nunes et al., 2014). Mobility may also play a significant role in the spread of arboviruses at more local scales; for example, time to travel has been shown to be strongly associated with incidence within distinct neighbourhoods of the same city during an arboviral outbreak (Faria, Lourenço et al., 2016). As real-time collection of genetic data becomes more frequent and available, increased time and spatial resolution will allow future phylogenetic and modelling studies to further evaluate the relative contribution of short- and long-distance mobility.

It may seem that these recently established approaches and real-time sequencing efforts are more appropriate for emerging viruses, such as ZIKV or the Ebola virus (Faria et al., 2017; Woolhouse, Rambaut, & Kellam, 2015) and that opportunities have been missed in the context of DENV, given its generalized geographical spread and endemicity. However, major gaps in our knowledge still exist in the context of DENV, sustained by the insufficiency of clinical reported case counts. In particular, how genotypes manage to persist in face of significant herd immunity, on the relative role of urban versus rural spread routes, off-season persistence and on the invasion and emergence of pathogenic lineages for which the displacement of the Asian/American by the Asian-1 type in South-East Asia is a good example (Lourenço & Recker, 2010; Hang, Holmes, Duong, Nguyen, & Tran, 2010). Hence, while the endemic state of DENV may render efforts of real-time sequencing less appealing, we argue that the enumerated knowledge gaps offer major research opportunities. It is also the case that retrospective analysis of archived clinical samples, such as blood banks, could contribute massively. However, real-time sequencing not only offers the same opportunities as retrospective approaches, as it also renders timely interventions possible with significant public health impact, for instance in face of emerging virulent genotypes or introduction of serotypes which may increase the number of heterologous challenges.

2.2 | Phylodynamic modelling frameworks

The genetic diversity of RNA viruses is shaped in direct response to changes in host and vector demographic and ecological factors, such as population size, structure or movement. This provides measurable connections between host population dynamics, viral evolution and transmission. In recent years, genetic, immunological and epidemiological data have been successfully used to investigate such connections, unifying key observations across biological scales within a single framework known as phylodynamics (Grenfell et al., 2004; Volz, Koelle, & Bedford, 2013). However, while some studies have

addressed dengue's epidemiology in the context of a phylodynamic framework (Rasmussen, Boni, & Koelle, 2014; Ke, Aaskov, Holmes, & Lloyd-Smith, 2013), integrative modelling approaches in which transmission models are used to simulate both the epidemiology and molecular evolution of the virus have not yet been applied to dengue.

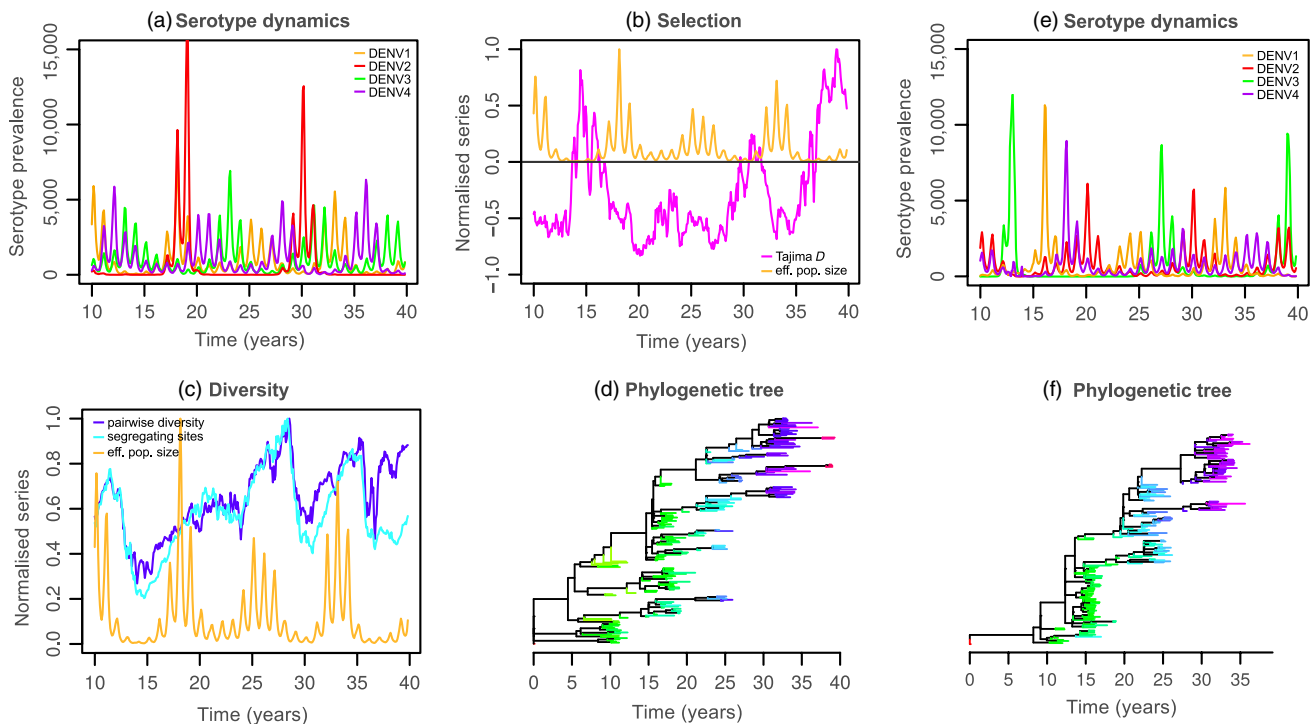
Dengue virus is a surprisingly conserved RNA virus (Choudhury, Lott, Banu, Cheng, & Teo, 2015; Lequime, Fontaine, Ar Gouilh, Moltini-Conclois, & Lambrechts, 2016; Thai, Henn, Zody, Tricou, & Nguyet, 2012; Amarilla, de Almeida, Jorge, Alfonso, & de Castro-Jorge, 2009; Holmes, 2003). Even though its serotypes present seemingly competitive dynamics, similar to the behaviour of antigenic strains of other fast evolving pathogens, observations of positive selection are rare (Tang et al., 2010; Filomatori et al., 2017; Waman et al., 2016; Hang et al., 2010; Choudhury et al., 2015; Martin, Chirivella, Co, Santiago, & Gubler, 2016; Sim, Aw, Wilm, Teoh, & Hue, 2015). Dengue transmission models have offered insight, but have so far failed to reach consensus on the possible biological drivers of the observed dynamic behaviour. For other pathogens, such as influenza A, phylodynamic transmission models have allowed to filter out a wide range of hypotheses regarding potential drivers (e.g., Bedford, Suchard, Lemey, Dudas, & Gregory, 2014; Minayev & Ferguson, 2009), as model validation becomes stricter and requires both epidemiological and evolutionary trajectories to match observed data patterns.

Here, we argue that phylodynamic models could offer critical insight into currently open questions on dengue's evolution and epidemiology, for example by assessing what assumptions about serotype immune interactions are compatible with empirical population genetic and phylogenetic data, similar to approaches applied to influenza A. This approach is exemplified in Box I, where we extended a previously published, individual-based dengue transmission model (Lourenço & Recker, 2013) to include explicit molecular evolution. Two scenarios are compared: one in which the dynamics and evolutionary histories are purely driven by demographic and transmission processes without immune interactions (Box I, a–d), and one where we considered temporary, serotype-transcending immunity following recovery from infection (Box I, e–f). With this approach, the phylogenetic output would be particularly informative as cross-immunity results in more imbalanced phylogenetic trees, akin to a ladder-like topology that is often reported for other pathogens, such as influenza A (Bedford et al., 2014; Bedford, Cobey, & Pascual, 2011) or within-host HIV (Shankarappa, 1999), where cross-immunity is known to play a crucial role.

Given seasonal epidemic troughs in endemic regions, possible mechanisms for viral persistence are still open topics of discussion, in particular as phylogenetic studies have shown that the lack of case reports can be followed by resurgence of previously circulating genetic variants (Bennett et al., 2003, 2010; Carrington, Foster, Pybus, Bennett, & Holmes, 2005). Persistence may be achieved by a combination of temporary circulation in the vector population (i.e., vertical transmission) (Bosio, Thomas, Grimstad, & Rai, 1992; Adams & Boots, 2010), an unknown reservoir (Weaver & Vasilakis, 2009; Vasilakis, Cardosa, Hanley, Holmes, & Weaver, 2012), silent transmission (Kyle & Harris, 2008; Morrison et al., 2010; Poblaj, Nitatpattana, Chaimarin, Barbazan, & Chauvancy, 2006), or may simply be a consequence of

Box I Phylodynamic modelling

The explicit modelling of molecular evolution allows for an exploration of dengue's drivers for key observations of phylogenetics and population genetics, such as tree imbalance, Tajima D and measures of viral diversity. Here, in order to illustrate such potential, we expand an individual-based framework to include the transmission and evolution of individual virions. We make a number of simplifying assumptions: one virion is modelled per host; the transmission chain of a single serotype is used to define the discrete birth–death process of virions; mutation occurs over the birth–death process, resulting in a multichotomous, evolution tree; a fixed molecular clock is used for the mutational input, which follows an infinite-sites model with no recombination; all mutations are assumed to be neutral; a burn-in period of 25 years for transmission (ending at $t = 0$ in figures), followed by 10 years for molecular evolution ($t = 10$), is considered before model output is recorded; a single founder virus is used ($t = 0$); population genetic sampling is uniform in time- and frequency-dependent; phylogenetic tree is a small sample ($<1\%$) of the evolution tree.



(a, b, c, d) Model output without considering serotype-immune interactions. (e, f) Model output with 9 months of transcending cross-immunity. A particular serotype (DENV1, orange) is chosen for the population genetics and phylogenetic output. Effective population size is considered to be the total number of virions of the chosen serotype. Some output variables are normalized between 0 and 1 for visualization purposes. (a and e) Dynamics of the four serotypes. (b) Tajima D (magenta) and effective population size (orange). (c) Number of segregating sites (light blue), sequence pairwise diversity (dark blue) and effective population size (orange). (d and f) Phylogenetic tree with tips coloured according to accumulated number of mutations (tips are sampled after the 10 year burn-in).

These figures present the dynamic potential of an individual-based model with explicit molecular evolution. In figures a–d, the complete model output shows how demographic stochasticity and the mechanics of transmission alone can dictate the intrinsic behaviour of viral diversity and phylogenetic shape. In contrast, figures e and f show how such signatures (e.g. phylogenetic shape) can be significantly altered when changing model assumptions—in this particular case, the addition of a 9-month period of transcending, cross-protective immunity.

reintroduction from nearby regions following local extinction. The particular hypothesis of continuous silent transmission is critical for public health, while the possibility of exogenous circulation has severe implications for control. Box I shows that viral diversity can be addressed in phylodynamic models, which is seen to respond both to seasonal troughs and to natural fluctuations in serotype prevalence. Diversity

can be expected to behave differently, for example depending on whether exogenous reservoirs are considered or not. Phylodynamic models could thus be used to explore hypotheses on mechanisms of (local) viral persistence in ways not possible with current transmission models and may therefore change our current understanding of the persistence of dengue viruses.

In a broader evolutionary perspective, drivers of key observations on the molecular evolution of dengue could also be studied within these frameworks. For instance, strong purifying selection is consistently reported in sequence data, for which it is generally accepted that the two-host life cycle imposes strong evolutionary constraints. Box I (b) shows, however, that even without explicit constraints on molecular evolution (e.g., deleterious mutations), negative Tajima D, a proxy for purifying selection, is self-emergent. As such, demographic and transmission processes appear sufficient to maintain the viral population as under purifying selection. Sensitivity experiments could therefore be used to test which processes seem to dictate these population genetic signatures; for example, how does host mixing in terms of mobility or patch organization alter such observations?

There are also reports of sporadic episodes of positive selection and fixation of particular lineages (Tang et al., 2010; Filomatori et al., 2017; Waman et al., 2016; Hang et al., 2010; Choudhury et al., 2015; Martin et al., 2016), and it is generally believed that most events of lineage replacement go unnoticed, as the majority of endemic regions rely on syndromic surveillance, and as these events are not expected to result in significant changes in disease incidence (Lourenço & Recker, 2010). While there are rare examples of well-described lineage shifts in the literature (OhAinle et al., 2011; Bennett, Holmes, Chirivella, Rodriguez, & Beltran, 2006), not much is known about the epidemiological and ecological contexts in which emergence of fitter lineages is favoured. Our previous work has suggested that stochastic effects and clonal interference are major determinants of the emergence of fitter, potentially more virulent variants (Lourenço & Recker, 2010). With these more detailed individual-based model frameworks, the strength of such determinants could be tested with higher resolution, discerning, for instance, between stochastic pressures dictated by population structure, host mixing or spatial heterogeneity in herd immunity (Lourenço & Recker, 2013).

Disparate levels of within- and between-host genetic diversity are a common feature of DENV viruses, and complementary hypotheses exist for these observations (Choudhury et al., 2015; Lequime et al., 2016; Thai et al., 2012). By allowing parameterization of mutational input and quantification on how population-level events modulate within-host genetic diversity, phylodynamic models could help make predictions on the relative contribution of different mechanisms. In particular, transmission bottlenecks could be implemented and their effect tested in the background of cross-immunity and clonal interference (Lourenço & Recker, 2010).

There are concerns related to viral escape variants following intervention programmes based on vaccination and/or Wolbachia-infected mosquitoes. While for Wolbachia-based control strategies, it can be argued that these concerns are so far theoretical (Bull & Turelli, 2013), escape in a postvaccination era has been reported for a wide range of pathogens (Read, Baigent, Powers, Kgosana, & Blackwell, 2015; Pérez-Sautu, Costafreda, Caylà, Tortajada, & Lite, 2011). The explicit generation, emergence and spread of escape mutants could be modelled within phylodynamic frameworks such as the one presented in Box I, given that individual mutations are already tracked. Vaccination, as demonstrated in previous work, can also be easily included (Lourenço

& Recker, 2016; Flasche et al., 2016). Parameterization on the efficiency of viral escape to vaccine-induced immunity would allow for sensitivity experiments and projections on the public health impact of such escape variants. Furthermore, demographic and ecological heterogeneities could also be researched, such as population structure, in order to evaluate under which conditions such variants could more easily emerge and devise appropriate intervention schemes that could stop their fixation.

Finally, as described in the previous section, real-time collection and analysis of genetic data will soon offer vast amounts of high-resolution data on population genetic measurements and phylogenetics, both in time and space. This will fundamentally change our perspective and the number of empirical observations that can be used to validate model frameworks. The use of more complex modelling approaches, such as individual-based models, including atypical dimensions such as detailed spatial scales (Box II) or molecular evolution (Box I), will not only be a valuable path of research, but could indeed become necessary approaches to be able to match the rich variety of observed data patterns across biological scales.

2.3 | Within-host models

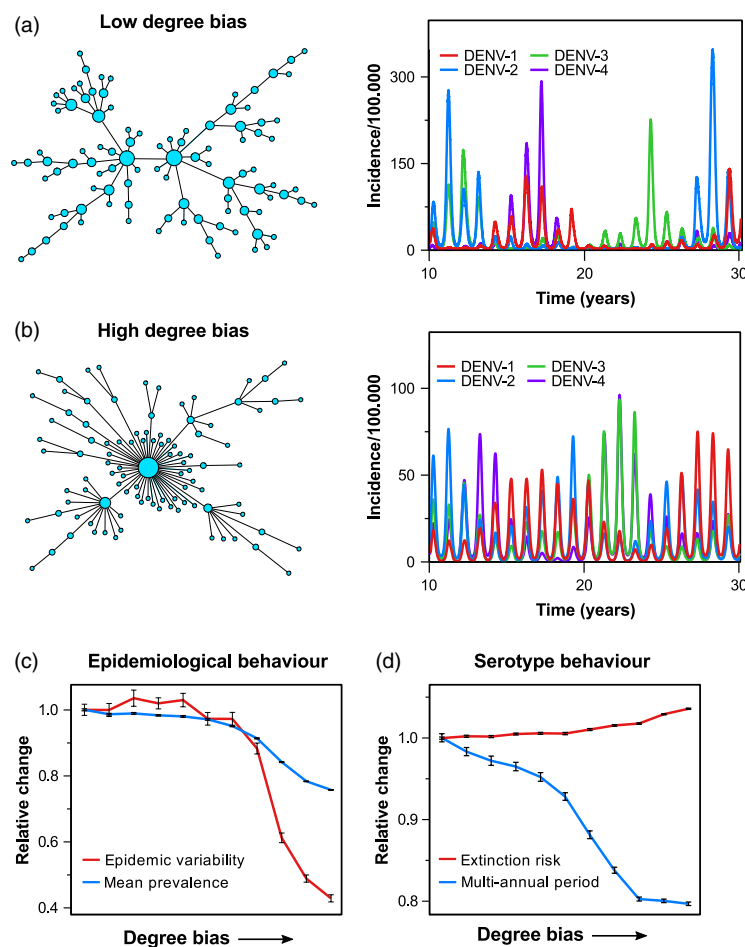
The vast majority of dengue models has focused on viral dynamics at the population-level. Given the importance that the within-host processes (e.g., ADE and cross-immunity) play for both viral transmission and disease pathology, it is thus surprising that just over a handful of studies have so far investigated viral dynamics at the within-host level (Nuraini, Tasman, Soewono, & Sidarto, 2009; Ansari & Hesaaraki, 2012; Clapham, Tricou, N. V., Simmons, & Ferguson, 2014; Ben-Shachar & Koelle, 2014; Gujarati & Ambika, 2014; Nikin-Beers & Ciupe, 2015; Clapham, Quyen, Kien, Dorigatti, & Simmons, 2016; Ben-Shachar, Schmidler, & Koelle, 2016). This could, however, be explained by the little understanding we currently have about the interactions between the virus, its target cells and the host immune responses that eventually clear infection, which may also be directly responsible for clinical outcome.

The target cell population for DENV replication is still not well characterized in vivo, although in vitro studies strongly suggest that monocytes, dendritic, endothelial and epithelial cells are among the preferential targets of infection (Fernandez-Garcia et al., 2009; Ubol & Halstead, 2010; Dejnirattisai et al., 2010; Katzelnick, Coloma, & Harris, 2017). Similarly, it has been demonstrated that both innate and adaptive immune responses play a role during infection, but it is not well understood how these can mechanistically and synergistically achieve viral clearance.

There are two main known mechanisms of antibody action for viral clearance in dengue infection—antibodies can indirectly act to clear infected cells via antibody-dependent cell-mediated cytotoxicity (ADCC), or directly by neutralization. A data-driven study by Clapham et al. (2016) focused on the distinction between these two mechanisms and found that either could explain the data (viral and antibody titres) equally well. Two other studies have focused on the general mechanics of humoral responses (B cell and free antibody) using different

Box II Sensitivity analysis using GPU acceleration

The increased computational power due to GPU parallelization permits a deeper exploration into the epidemiological effects of different community structures, vector and host heterogeneities, environmental heterogeneities or human movement patterns. Here, in order to illustrate the use of GPU-accelerated agent-based modelling, we investigate how the connectivity between subpopulations, as a proxy for human movement/major commuting patterns, affects the epidemiological dynamics of dengue within a spatial meta-population model. In this framework, the population (human and mosquitoes) is divided into subpopulations and arranged spatially into a network of communities, connected by major trade and/or commuting routes. Dengue transmission events are dispersed through the network predominantly via local connections between communities as well as occasional but random transmission events between distant (i.e., not directly connected) subpopulations. The global connectivity pattern can then be quantified by means of the network's *degree bias*, which, in its simplest form, can be interpreted as the number of nodes (i.e., communities) that are only connected to one other node. Plots a–b show two examples of networks with either a low or a high degree bias, respectively. As can be seen in the corresponding time series of serotype-specific incidence, the way that communities are connected to form a meta-population has a strong effect on the epidemiological dynamics, especially with respect to the interannual variability and serotype periodicity. This can be further illustrated by means of a sensitivity analysis, where we gradually increased the degree bias and recorded the model's relative change in four characteristic epidemiological and serotype-specific measures (plots c–d). In this specific example, where we considered a population size of 5 million humans and up to 6 million mosquitoes, we gained a speed-up of around 320 (using a Nvidia GTX 1070 graphics card). In other words, running the same analysis, which took just over 9 hr to complete, would have taken about 4 months on a single core CPU and still many weeks using multithreaded CPU computing on a high-end PC. This clearly illustrates how GPU-based computing may facilitate the use of bigger and more complex models for addressing some of the outstanding dengue epidemiological and evolutionary questions discussed in this work.



(a, b) The degree of connectivity between subpopulations has a strong effect on the epidemiological dynamics of dengue. (c, d) Model sensitivity analysis showing the relative change in epidemiological and serotype-specific characteristics depending on the network's degree bias. Each point is the result of 50 model runs over a 100-year period, with error bars depicting the standard error.

modelling frameworks (Gujarati & Ambika, 2014; Nikin-Beers & Ciupe, 2015). While one study concluded that enhancing antibodies (ADE) were key in secondary infections (Gujarati & Ambika, 2014), the other study suggested instead a direct decrease of overall heterologous viral clearance, whereby cross-reactive antibodies render virions unavailable for further binding and subsequent clearance by ADCC (Nikin-Beers & Ciupe, 2015). In another study, the question about the role of the adaptive and innate immune responses in disease severity was addressed (Ben-Shachar & Koelle, 2014; Ben-Shachar et al., 2016). It was shown that characteristic features of primary infection could be replicated solely by innate immune responses (NK cells). In contrast, features of secondary infection, including ones related to clinical outcome, required greater infectivity rates (ADE) together with T cell-mediated clearance of infected cells in a process arguably similar to well-described cytokine storms (Rothman, 2011; Katzelnick et al., 2017). In line with previous results by Clapham et al. (2014), it was proposed that the observed variations in viral load could largely be explained by assuming patient-specific incubation periods.

Generally and similarly to the aforementioned discrepancy between model assumptions related to immune interactions at the population level, within-host approaches have been able to replicate important and ubiquitous data patterns, but, either due to formalism or data restrictions, have been limited in establishing a consensus framework. In some cases, the models and the data sets differed, making it hard to assess whether contrasting conclusions are data- or model-driven. For instance, some models propose a key role of innate immunity in clearance of primary infection (Ben-Shachar & Koelle, 2014; Ben-Shachar et al., 2016), while others propose that either humoral or cell-mediated clearance is required (Nuraini et al., 2009; Ansari & Hesaaraki, 2012; Clapham et al., 2014; Gujarati & Ambika, 2014; Nikin-Beers & Ciupe, 2015; Clapham et al., 2016). Equally, asymmetries between serotypes in both cell infectivity and timing of peak viraemia versus symptomatic manifestations were reported in one study (Ben-Shachar et al., 2016) but not in another (Clapham et al., 2014).

A current major problem is the lack of fine-scaled data that captures the early, prepeak infection dynamics. This issue arises from the self-limiting nature of dengue infections, fast viral clearance rates and the absence of antiviral therapies against dengue, which effectively prevent human infection studies. In fact, antiviral therapy has been devalued due to the fact that the timing of clinical symptoms often overlaps with peak viral load (Clapham et al., 2014); that is, most patients will seek treatment at a point when the virus is already being cleared by the immune system. However, in another example of contrasting observations, if there are indeed significant differences between the serotypes, as suggested by Ben-Shachar et al. (Ben-Shachar et al., 2016), then there might be a question whether sero-specific strategies could be beneficial, for which modelling studies could contribute useful answers.

Another promising avenue is in the context of vaccination. Recently, the first dengue vaccine was licensed (Dengvaxia®) and many others are in advanced trial stages (Schmitz, Roehrig, Barrett, & Hombach, 2011; Thomas & Endy, 2011; Schwartz, Halloran, Durbin, & Longini, 2015). For a good number of these, data will be made

available on population-level efficacy but also on within-host markers and dynamic measurements of infection and immune response (e.g., Dorigatti, Aguas, Donnelly, Guy, & Coudeville, 2015). Such data can be used to inform new modelling frameworks to explain vaccine action (e.g., infection versus transmission blocking), which may have considerable impact on licensing and future distribution. We note, however, that is yet unknown whether fine-scale kinetic data will be available, which is essential to parameterize models (Magnus, 2013) and to add insight into current knowledge gaps on the fast kinetics of the virus, in particular during the initial phase of infection. Finally, and similar to between-host models, within-host models of dengue have so far only considered a single ecological scale. With the advent of models with better computational implementation (see next section), both the between-host and within-host dynamics could be integrated in unified, multiscale frameworks. These could be used to investigate how assumptions about viral dynamics and immune kinetics, together with individual-level variations and transmission mechanics, would influence epidemiological and evolutionary observations at the population level (Box I).

2.4 | GPU modelling frameworks

Many traditional approaches in theoretical epidemiology of dengue and other infectious diseases rely on mass-action principles whereby the rate at which a disease spreads through a population is directly proportional to either the number or the proportion of infected and susceptible individuals within that population. Under these assumptions, every individual has the same probability of getting infected, contributes equally to disease transmission and recovers at the same rate as everyone else. Their low computational footprint and analytical tractability make these models an attractive choice for investigating population-level dynamical behaviours, especially when homogeneity in time and space can safely be assumed. In many cases, this cannot be guaranteed, however, and individual variations, arising through the stochastic nature of infection events, for example, or through ecological and demographic heterogeneities, cannot be captured by these models and require a different approach altogether.

Agent or individual-based models offer a natural and intuitive way to account for these variations by keeping track of individual, probabilistic infection events in both humans and mosquito vectors and have been implemented to various degrees of realism and model complexity. As an example, spatial details can be included by dividing the population into smaller subpopulations arranged in a rectangular grid (see Figure 3). This approach has already highlighted how the spatial segregation of individuals, causing stochastic, local extinction and re-invasion, can explain much of dengue's observed epidemiology (Lourenço & Recker, 2013). Further detail may be introduced by adding more realistic spatial arrangements by means of complex networks with nodes representing villages or cities and edges representing their connecting trade or commuting routes, for example. Subpopulations can further be divided to take into account individual households, work places or schools, as well as the human movement patterns between them (see, e.g., (Chao, Halstead, Halloran, & Longini,

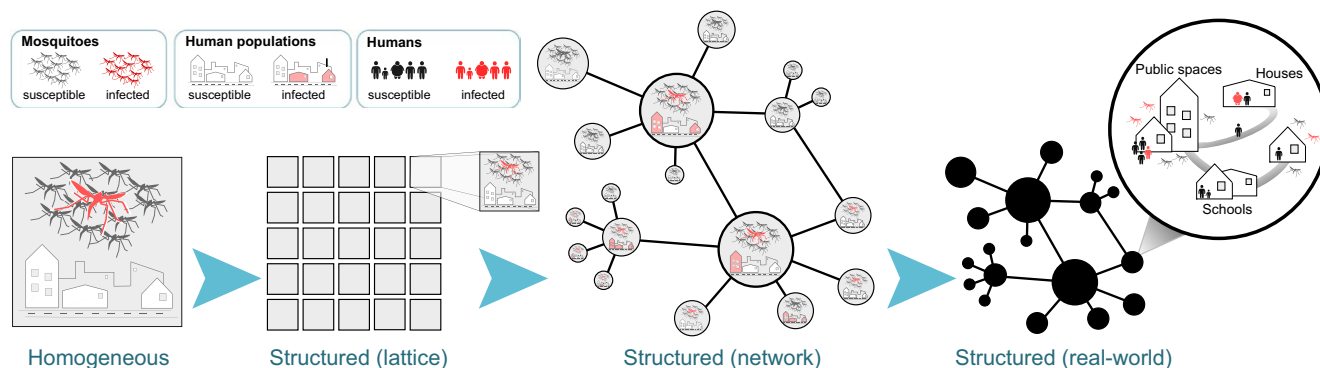


FIGURE 3 Increasing model complexity demands higher computational power. Model detail can be added by dividing a well-mixed population into separate subpopulations, arranged in a regular spatial grid or by means of complex networks to represent geographical distribution of villages, towns and cities, with edges corresponding to major human movement patterns. Depending on data availability, more spatial and demographic detail can be added by considering individual households, places of work or schools. However, the computational demands increase significantly with more detailed information to keep track of, making the model very setting-specific and impractical for sensitivity analyses and model fitting exercises

2012; Karl, Halder, Kelso, Ritchie, & Milne, 2014; Perkins, Garcia, Paz-Soldán, Stoddard, & Reiner, 2014)). In fact, the explicit description of each individual (human and vector), together with their ecological and socio-demographic interactions, allows a near limitless level of spatio-temporal detail to be incorporated into these models.

The inclusion of (every) minute detail comes at the cost of computational feasibility and generalizability, however. That is, the computational demands imposed by an increase in model complexity, due to the incorporation of more, and more detailed information of individual-level behaviours, can quickly exceed the capabilities of modern-day personal computers and require either very long run-times or the implementation on high-performance computer clusters. Furthermore, the limited availability of fine-scaled data necessary to parameterize these models often restricts their use to the investigation of particular aspects of dengue epidemiology and mostly in one specific setting. Results obtained from these studies, although highly informative for the particular research question, are therefore not easily transferable to other epidemiological or geographical settings. This then implies that for research questions of a more general nature a balance needs to be struck between a model's biological and ecological realism and computational feasibility.

In many cases, the computational overhead of agent-based models arises not necessarily from individual infection events but rather from keeping track of host and vector demography and ecology, for example the processes related to birth, death and ageing. As these are largely independent events, they are amenable to code parallelization and can thus benefit from graphics processing unit (GPU) acceleration, which specializes in attaining high arithmetic throughputs. Speed-ups of anywhere between 10 and 1000 times from a serialized central processing unit (CPU) code are theoretically feasible, although in practice this very much depends on the particular model and implementation. That is, the benefits of GPU acceleration can easily get lost on increasingly refined spatial resolutions, for example, when more time is spent keeping track of individuals' movements (which is generally not

parallelizable) than updating infection statuses or age. Nevertheless, the gain in computational speed-up through GPU acceleration enables us to simulate much bigger and more detailed models in significantly less time (see Box II).

The necessity to include higher-resolution in biological scales such as population structure in modelling frameworks is clear, given the recent rise of ever more detailed genetic (Faria et al., 2017; Woolhouse et al., 2015), mobility (Kraemer, Perkins et al., 2015; Wesolowski et al., 2015; Lemey et al., 2014) or social/communication (Salathé, Bengtsson, Bodnar, Brewer, & Brownstein, 2012) data sets. That is, the increased difficulties (e.g., interpretation) arising from the use of more complex models were not compensated for by sufficient data to validate model output; however, this is now changing. Methodologies such as the one presented in Box II now offer major opportunities to test and create new hypotheses on the spread, evolution and control of communicable diseases.

The spatial resolution of the research question dictates the complexity level of community structuring required in a model. Nonspatial deterministic approaches provide a natural entry point in understanding the epidemiology of pathogens strictly across time (Nagao & Koelle, 2008; Rodriguez-Barraquer, Mier-y Teran-Romero, Schwartz, Burke, & Cummings, 2014; Ferguson, Rodriguez-Barraquer, Dorigatti, Mier-y Teran-Romero, & Laydon, 2016). Epidemiological questions directed over very small geographical areas, such as investigating the effects of vector distribution, or vector and host movement between an individual's home and workplace, require fine-scale spatial models (Chao, Longini, & Halloran, 2013; Reiner, Stoddard, & Scott, 2014; Perkins, Paz-Soldan, Stoddard, Morrison, & Forshey, 2016; Hladish, Pearson, Chao, Rojas, & Recchia, 2016). Coarser-scaled network models are applied when individual movement between households becomes redundant, for example exploring the effects of national or international human movement, or environmental heterogeneities over large spatial regions, on the spread, evolution and control of communicable diseases.

In the context of DENV, such approaches are now reaching the capacity to be modulated and fitted to particular, existing host

populations and to simulate real-world scenarios in terms of transmission and control. This would be particularly valuable in the context of DENV vaccination modelling, which is recent in the literature (e.g., (Lourenço & Recker, 2016; Flasche et al., 2016; Chao et al., 2012; Coudeville & Garnett, 2012)). For most of the published studies, simplifying assumptions are made on population heterogeneities or age-dependent factors, for instance. However, with the amelioration of computational costs, realistic vaccination campaigns can be assessed, including details related to relative targeting of rural versus urban settings or detailed catch-up and routine strategies dependent on host age. In terms of vector control programmes, the actual location of water reservoirs, the influence of climate or host mobility networks can also be considered. In summary, these advanced computing techniques have the capacity to change our perspective of modelling frameworks from being based on groups, generally of individuals sharing similar properties or phenotypes, to a perspective of host-pathogen systems focusing on the role of single individuals, and how the mechanics of transmission and evolution at that scale can emerge as self-evident patterns at the level of the population.

3 | CONCLUDING REMARKS

We have reviewed the biological and epidemiological background of dengue, together with the major achievements of computational approaches dedicated to the pathogen. In addition and in the context of its increasing success as a global threat to public health, we have also highlighted critical knowledge gaps and research underachievements that call for an urgent renewed focus and promotion of such methodologies. We argue that possible advancements, based on new processing strategies, real-time data sources and modelling frameworks already implemented for other pathogens, are already at reach of the research community. These new approaches are expected to make a significant contribution to our understanding of the evolutionary ecology and immunology of the dengue virus and its control in the near future.

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CONFLICT OF INTEREST

The authors declare no conflict of interest.

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APPENDIX

Data Description

Brazilian data are for suspected case counts from the Sistema de Informação de Agravos de Notificação (national notifiable diseases information system, SINAN) database (Brazil SINAN, <http://portalsinan.saude.gov.br/>). The regions of north and south, as presented in Figure 1, present aggregated data (south: states of Paraná, Santa Catarina, Rio Grande do Sul; north: states of Rondônia, Acre, Amazonas, Roraima, Pará, Amapá, Tocantins).

Dengue incidence data for Puerto Rico were provided by Michael Johansson from the CDC Dengue Branch, comprising clinically suspected cases of dengue fever (DF) and dengue haemorrhagic fever (DHF) in Puerto Rico between 1986 and 2012. Serotype-specific testing has varied over time, so we adjusted the serotype-specific incidence data proportionally to match the incidence of all suspected cases by month. A shorter time series had previously been published and analysed in (Johansson, Cummings, & Glass, 2009).

Annual incidence of DHF in Thailand 1973–2009 is of clinically confirmed cases and was obtained from the annual epidemiological surveillance reports published by the Ministry of Public Health (website: epid.moph.go.th). Dengue serotype prevalence for Thailand 1973–1999 was based on children hospitalized at the Queen Sirikit National Institute of Child Health in Bangkok, as previously published in (Nisalak et al., 2003).

Yearly counts of dengue publications (1970–2016) were collected from PubMed under two lists (queries). The first contained the total number of publications per year dedicated to dengue, including all hits with reference to dengue in the title and abstract (counts collected using the “results by year” functionality in PubMed). The second was a subset of the first, which included the use of dynamic models in the context of dengue research (manually curated). Queries were performed on the 1st of February 2017.